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[wherein: R1 = -(CH₂)₁₋₂-S(O)₀₋₂-(C₁₋₆ alkyl) or (un)substituted (cyclo)alkyl, alk(en/yn)yl, (hetero)aryl, etc.; R2 = H, C₁₋₆ alkyl optionally substituted with 1-3 substituents, (CH₂)₀₋₄-(hetero)aryl, C₂₋₆ alk(en/yn)yl, etc.; R3 = H, C₁₋₆ alkyl optionally substituted with 1-3 substituents, (CH₂)₀₋₄-(hetero)aryl, etc.; R4 = C₁₋₁₀ alkyl optionally substituted with 1-3 substituents, -(CH₂)₀₋₃cycloalkyl, -(CR₇R₈)₀₋₄-(hetero)aryl, etc.; one of R₅ and R₆ is H and the other is -C(O)(CR₉R₁₀)₁₋₆-X-R₁₁, etc.; R₇ and R₈ are independently selected from H, alkyl, hydroxyalkyl, alk(en/yn)yl, etc.; R₉ and R₁₀ are independently selected from H or C₁₋₁₀ alkyl; R₁₁ = (hetero)aryl, optionally substituted C₁₋₁₀ alkyl, or C₃₋₈ cycloalkyl, etc.; X = O, S, SO₂, etc.]. Compds. I include inhibitors of beta-secretase enzyme useful in the treatment of Alzheimer's disease and other diseases characterized by deposition of A beta-peptide in a mammal. Biol. examples include beta-secretase inhibition, assays using synthetic oligopeptide-substrates, inhibition of A beta production in human patients, etc. For instance, compound II (preparation 8)

was prepared via amidation of benzoic acid derivative III by diamino(hydroxy)propane derivative IV and subsequent Boc-cleavage (no yield data). Using ¹⁹F-NMR an intramol. acyl-migration was observed when compound II was dissolved in DMSO-¹⁹F and pH 4 buffer solution was added.

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